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STATEMENT BY APPLICANT**
(Not for submission under 37 CFR 1.99)

Application Number	10568051
Filing Date	2006-02-10
First Named Inventor	Miller
Art Unit	3743
Examiner Name	(not yet assigned)
Attorney Docket Number	PR60436USw

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	2	5861473		1999-01-19	DECROSTA et al.		
	3	6068789		2000-05-30	BARNES et al.		
	4	6241828		2001-06-05	BARNES et al.		

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	1	20050092679	A1	2005-05-05	WARBY		

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Receipt date: 01/18/2008

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	1	0134847	EP	A1	1985-03-27	Trutek Research Inc.	<input type="checkbox"/>
	2	9312161	WO	A1	1993-06-24	Schering Corporation	<input type="checkbox"/>
	3	02072448	WO	A1	2002-09-19	Chiesi Farmaceutici s.p.a.	<input type="checkbox"/>
	4	03049786	WO	A2	2003-06-19	Glaxo Group Limited	<input type="checkbox"/>

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	1	ITFG/IPAC-RS COLLABORATION - CMC LEACHABLES AND EXTRACTABLES TECHNICAL TEAM; "Leachables and Extractables Testing: Points to Consider", A Response to the FDA draft Guidance for Industry; March 27, 2001; pp. 1-36; < www.fda.gov/ohrms/dockets/ac/00/reports/3657_rpt1.pdf >	<input type="checkbox"/>
	2	JOSEPH H. GROEGER & LESLIE M. COMPTON; "Identifying and Preventing Contamination from Pharmaceutical Packaging"; Medical Plastics and Biomaterials Magazine; May 1997; < www.devicelink.com/mpb/archive/97/05/005.html >	<input type="checkbox"/>
	3	U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES - FOOD AND DRUG ADMINISTRATION - CDER; "Draft Guidance for Industry - Metered Dose Inhaler (MDI) and Dry Powder Inhaler (DPI) Drug Products"; Chemistry, Manufacturing, and Controls Documentation; October 1998; pp. 1-62; < www.fda.gov/Cder/guidance/2180dft.pdf >	<input type="checkbox"/>

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